

~~R₂ is C₁₋₆alkyl;~~

~~R₃ is selected from hydrogen, C₁₋₆alkyl, mono- or di(C₃₋₆cycloalkyl)methyl, C₃₋₆cycloalkyl; C₃₋₆alkenyl; hydroxyC₁₋₆alkyl, C₁₋₆alkylcarbonyloxyC₁₋₆alkyl and C₁₋₆alkyloxyC₁₋₆alkyl;~~

~~R₄ and R₅ are independently selected from C₁₋₈alkyl, mono- or di(C₃₋₆cycloalkyl)methyl, Ar¹CH₂, C₃₋₆alkenyl, C₁₋₆alkyloxyC₁₋₆alkyl, hydroxyC₁₋₆alkyl, thienylmethyl, furanymethyl, C₁₋₆alkylthioC₁₋₆alkyl, morpholinyl, mono- or di(C₁₋₆alkyl)aminoC₁₋₆alkyl, di(C₁₋₆alkyl)amino, C₁₋₆alkylcarbonylC₁₋₆alkyl, C₁₋₆alkyl substituted with imidazolyl; or a radical of the formula -(C₁₋₆alkanediyl)-O-CO-Ar¹;~~

~~or R₃ and R₄ taken together with the nitrogen atom to which they are attached form a pyrrolidinyl, piperidinyl, homopiperidinyl or morpholinyl group, optionally substituted with C₁₋₆alkyl or C₁₋₆alkyloxy;~~

~~Ar is selected from phenyl substituted with 1, 2 or 3 substituents independently selected from halo, C₁₋₆alkyl, trifluoromethyl, cyano, C₁₋₆alkyloxy, benzyloxy, C₁₋₆alkylthio, nitro, amino and mono- and di(C₁₋₆alkyl)amino; and pyridinyl substituted with 1, 2 or 3 substituents independently selected from halo, C₁₋₆alkyl, trifluoromethyl, hydroxy, cyano, C₁₋₆alkyloxy, benzyloxy, C₁₋₆alkylthio, nitro, amino, mono- and di(C₁₋₆alkyl)amino and piperidinyl; and~~

~~Ar¹ is selected from phenyl, pyridinyl, and phenyl substituted with 1, 2 or 3 substituents independently selected from halo, C₁₋₆alkyl, C₁₋₆alkyloxy, di(C₁₋₆alkyl)aminoC₁₋₆alkyl, trifluoromethyl and C₁₋₆alkyl substituted with morpholinyl.~~

Please add the following claims:

19. (new) The method of claim 5 wherein said disorder is anxiety or depression.

20. (new) The method of claim 5 wherein said disorder is irritable bowel syndrome, inflammation, anorexia nervosa, seizures, or substance abuse.